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Effect of haemoglobin levels on outcome in intravenous thrombolysis-treated stroke patients

Thrombolysis Stroke Patients TriSP

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Abstract

Introduction: Alterations in haemoglobin levels are frequent in stroke patients. The prognostic meaning of anaemia and polyglobulia on outcomes in patients treated with intravenous thrombolysis is ambiguous.

Patients and methods: In this prospective multicentre, intravenous thrombolysis register-based study, we compared haemoglobin levels on hospital admission with three-month poor outcome (modified Rankin Scale 3–6), mortality and symptomatic intracranial haemorrhage (European Cooperative Acute Stroke Study II-criteria (ECASS-II-criteria)). Haemoglobin level was used as continuous and categorical variable distinguishing anaemia (female: <12 g/dl; male: <13 g/dl) and polyglobulia (female: >15.5 g/dl; male: >17 g/dl). Anaemia was subdivided into mild and moderate/severe (female/male: <11 g/dl). Normal haemoglobin level (female: 12.0–15.5 g/dl, male: 13.0–17.0 g/dl) served as reference group. Unadjusted and adjusted odds ratios with 95% confidence intervals were calculated with logistic regression models.

Results: Among 6866 intravenous thrombolysis-treated stroke patients, 5448 (79.3%) had normal haemoglobin level, 1232 (17.9%) anaemia – of those 903 (13.2%) had mild and 329 (4.8%) moderate/severe anaemia – and 186 (2.7%) polyglobulia. Anaemia was associated with poor outcome (OR_{adjusted} 1.25 (1.05–1.48)) and mortality (OR_{adjusted} 1.58 (1.27–1.95)). In anaemia subgroups, both mild and moderate/severe anaemia independently predicted poor outcome

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(OR_{adjusted} 1.29 (1.07–1.55) and 1.48 (1.09–2.02)) and mortality (OR_{adjusted} 1.45 (1.15–1.84) and OR_{adjusted} 2.00 (1.46–2.75)). Each haemoglobin level decrease by 1 g/dl independently increased the risk of poor outcome (OR_{adjusted} 1.07 (1.02–1.11)) and mortality (OR_{adjusted} 1.08 (1.02–1.15)). Anaemia was not associated with occurrence of symptomatic intracranial haemorrhage. Polyglobulia did not change any outcome.

Discussion: The more severe the anaemia, the higher the probability of poor outcome and death. Severe anaemia might be a target for interventions in hyperacute stroke.

Conclusion: Anaemia on admission, but not polyglobulia, is a strong and independent predictor of poor outcome and mortality in intravenous thrombolysis-treated stroke patients.

Keywords

Anaemia, polyglobulia, haemoglobin, intravenous thrombolysis, outcome, stroke

Date received: 19 July 2019; accepted: 21 October 2019

Introduction

Alterations in haemoglobin levels (HLs) on admission are frequently (anaemia up to 25%) observed in acute stroke patients.^{1,2} In general stroke populations, anaemia was associated with poorer outcomes^{1,3–5} except for one study.⁶ Only one smaller study (n = 217) has investigated the effect of anaemia on outcomes in stroke patients treated with intravenous thrombolysis (IVT). In this study, the development of anaemia or worsening of anaemia in the first days after admission was associated with poor functional outcome and mortality but not the presence of anaemia on admission.⁷ A second study, including both IVT (n = 466) and endovascular treated patients (n = 712), found anaemia on admission being an independent predictor of poor functional outcome and mortality.⁸ From a pathophysiological point of view, low HL at stroke onset is likely to impair outcomes due to a mismatch between increased metabolic requirements of the penumbral brain tissue and lowered oxygen transport capacity and reduced blood perfusion. Furthermore, HL might also affect outcomes via alterations in cerebrovascular autoregulation, blood coagulation and inflammatory mediators.^{1,9–12}

On the other side of the spectrum, the presence of polyglobulia in the general stroke population was associated with mortality in one¹ but not in another study.⁴ Therefore, the prognostic meaning of anaemia and polyglobulia on outcomes in patients treated with IVT remains unclear.

The aim of this study was to investigate the effect of baseline HL on functional outcome, mortality and bleed-risk in a large cohort of IVT-treated stroke patients.

Methods

For this cohort study, we used prospectively collected data from the *Thrombolysis in Ischemic Stroke*

Patients (TRISP) registry which has been previously described.¹³ Ten TRISP centres participated in this study (eTable 1). A complete list of all TRISP centres is presented in the online supplement (eAppendix 1). Data collection was done locally in each stroke centre using a standardised form with predefined parameters.^{14,15} Data of the local registries were pooled and analysed at the stroke centre Basel. Parameters of interest for the present study were age, sex, National Institutes of Health Stroke Scale (NIHSS) score,¹⁶ blood pressure prior to IVT treatment, onset-to-treatment time, estimated glomerular filtration rate using the CKD-EPI formula,¹⁷ glucose levels and HL in blood serum on admission, vascular risk factors according to predefined criteria^{13,18} and prior treatment with antithrombotic agents (antiplatelet agents or anticoagulants). Outcome parameters were mortality and the modified Rankin Scale (mRS) score at three months assessed either by outpatient visits or telephone calls with patients and/or relatives. Poor functional outcome was defined as a mRS score of 3–6. As safety outcome we defined the occurrence of symptomatic intracranial haemorrhage (sICH) using the ECASS-II-criteria.¹⁹ Intracranial haemorrhage was monitored by follow-up CT or MRI – in most cases performed at 24 h after start of IVT and immediately in case of neurological deterioration – as described in prior research.^{15,20}

Included data were collected up to 30 January 2014 (eTable 1). All patients with missing data on (i) HL, (ii) three-month outcome and (iii) sICH-data were excluded.

Statistical analyses

Statistical analyses were performed with SPSS Statistics version 25 (IBM) and STATA version 14.1 (StataCorp LP).

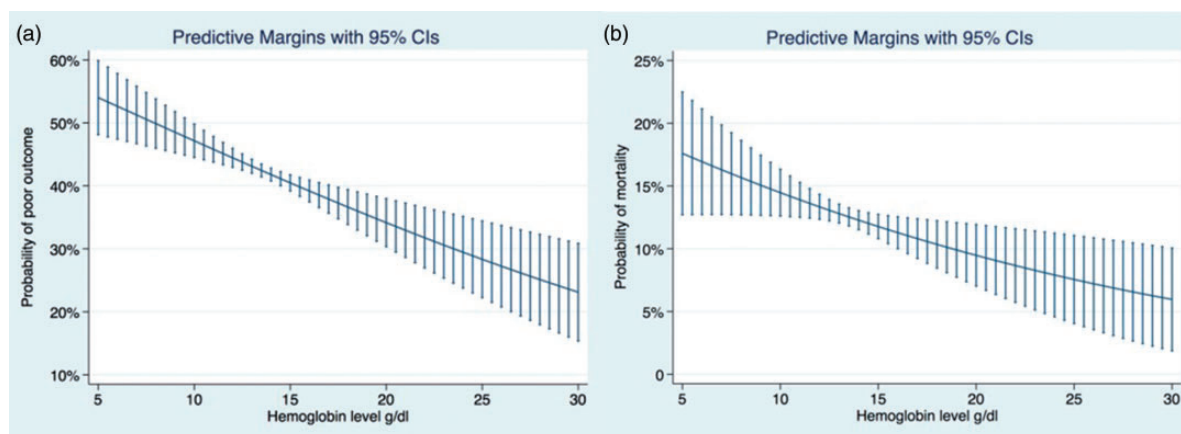


Figure 1. Predictive margins with 95% CI of the adjusted logistic regression models showing the association of haemoglobin levels and outcomes in IVT-treated stroke patients: (a) adjusted for age, stroke severity (NIHSS), glucose, eGFR, RF diabetes and (b) adjusted for age, stroke severity (NIHSS), RF diabetes.

We investigated associations between HL and outcome measures using HL as a (i) continuous variable and as a (ii) categorical variable distinguishing anaemia (female: <12 g/dl; male: <13 g/dl) and polyglobulia (female: >15.5 g/dl; male: >17 g/dl) defined by the criteria of the World Health Organisation (WHO).²¹ Normal HL (female: 12–15.5 g/dl; male: 13–17 g/dl) served as reference group. According to the grading system of the WHO, anaemia was further subdivided into mild (female: 11–11.9 g/dl; male: 11–12.9 g/dl) and moderate/severe (female/male: <11 g/dl).²²

Continuous data were summarised as median and interquartile range. We used Chi²-test and Fisher's exact test for categorical variables where appropriate and the Mann–Whitney U-test for continuous variables. The association between HL and each outcome was estimated by calculating odds ratios (OR) with 95% confidence intervals (95% CI), using binary logistic regression models. All variables with $p < 0.05$ were included in the multivariable analyses using stepwise regression with backward elimination. To avoid overfitting, the maximum number of potential confounders in the final model was restricted to one-tenth of the number of outcome events. Predictive margins of the adjusted logistic regression models are displayed in Figure 1.

Receiver operating characteristic (ROC) and area under the curve (AUC) were calculated to show the accuracy of HL to predict poor outcome and mortality.

The study was approved by the ethics committee in Basel, Switzerland. The requirement for additional local ethical approval differed between participating centres and was obtained if required. Anonymised data will be shared by request from any qualified investigator.

Results

Data were eligible for analysis in 6866 (92.8%) of the 7395 IVT-treated patients. Reasons for excluding patients were missing data on HL ($n = 213$; 2.9%), three-month outcome ($n = 257$; 3.5%) or sICH ($n = 59$; 0.8%).

Among study patients, 5448 (79.4%) had normal HL, 1232 (17.9%) anaemia and 186 (2.7%) polyglobulia.

Anaemia versus normal HL

Baseline characteristics are presented in Table 1. Patients with anaemia were older, had more severe strokes, lower median systolic blood pressure, lower eGFR at stroke onset, more frequently on antithrombotics (antiplatelets and/or anticoagulation) and were more likely to have a history of atrial fibrillation, coronary artery disease, diabetes mellitus, hypertension and prior stroke compared to patients with normal HL. Patients with anaemia more often had poor functional outcome (55.5% versus 39.5%) and died more often (22.4% versus 11.3%) during the three-month follow-up, while the proportion of patients with symptomatic ICH did not differ significantly (4.5% versus 4.3%). Data of recurrent strokes within three months after stroke onset were available in a subgroup of patients only ($n = 2518$, centres of Amsterdam, Basel, Belgrade, Bern, Brescia and Modena). Frequency of recurrent stroke did not differ significantly between patients with anaemia and normal HL (3.3% versus 3.4%) as well as between polyglobulia and normal HL (3.3% versus 4.9%). (Table 1)

Anaemia was associated with poor functional outcome ($OR_{unadjusted}$ 1.91, 95% CI 1.69–2.17 and $OR_{adjusted}$ 1.25, 95% CI 1.05–1.48) and mortality

Table 1. Clinical characteristics and frequency of outcome events of IVT-treated stroke patients divided into groups depending on their haemoglobin level (HL) at stroke onset.

	Normal HL n = 5448	Anaemia ^a n = 1232	Normal HL versus anaemia P value	Mild anaemia ^b n = 903	Normal HL versus mild anaemia P value	Moderate/ severe anaemia ^c n = 329	Normal HL versus moderate/ severe anaemia P value	Polyglobulia ^d n = 186	Normal HL versus polyglobulia P value
Hb on admission, in g/dl, median (IQR)	14.2 (13.4–14.9)	11.6 (10.9–12.2)	<0.001	11.9 (11.5–12.4)	<0.001	10.2 (9.5–10.6)	<0.001	16.7 (16.0–17.4)	<0.001
Age, years, median (IQR)	70 (59.8–78)	76 (67–82)	<0.001	76 (67–82)	<0.001	77 (66–83)	<0.001	64.9 (57–75)	0.001
Men, n (%)	3158 (58)	697 (56.6)	0.371	569 (63.0)	0.005	128 (38.9)	<0.001	78 (41.9)	<0.001
Stroke severity, NIHSS, median (IQR)	9 (5–15)	12 (7–17)	<0.001	11 (6–17)	<0.001	13 (8–18)	<0.001	9 (5–15)	0.870
Systolic blood pressure, mmHg, median (IQR)	157 (140–173)	150 (136–169)	<0.001	151 (138–170)	<0.001	150 (132–164)	<0.001	163 (148–180)	<0.001
Onset-to-treatment, min, median (IQR)	142 (105–180)	145 (110–180)	0.215	145 (109–180)	0.316	145 (110–185)	0.400	138 (95.5–184)	0.550
eGFR on admission ml/min, median (IQR)	80.6 (63.1–93)	72.2 (51.1–88)	<0.001	73.4 (53.5–88.1)	<0.001	68.8 (43.7–87.5)	<0.001	80.3 (64–92.7)	0.766
Glucose on admission mmol/l, median (IQR)	6.6 (5.8–7.9)	6.5 (5.6–7.9)	0.019	6.6 (5.7–8.0)	0.348	6.3 (5.4–7.8)	0.001	6.95 (6.0–8.88)	0.015
Prior antithrombotics, n (%)	1851 (35.1)	588 (48.9)	<0.001	419 (46.4)	<0.001	169 (51.4)	<0.001	62 (34.5)	0.696
Antiplatelets, n (%)	1504 (30.8)	466 (42.1)	<0.001	333 (40.8)	<0.001	133 (45.5)	<0.001	55 (30.6)	0.953
Anticoagulation with or without antiplatelets, n (%) ^e	195 (4.0)	70 (6.3)	0.001	54 (6.6)	0.001	16 (5.5)	0.211	7 (3.9)	0.946
Atrial fibrillation, n (%)	1388 (25.7)	409 (33.7)	<0.001	293 (32.4)	<0.001	116 (35.3)	<0.001	55 (29.6)	0.234
Hypertension, n (%)	3569 (65.6)	892 (72.6)	<0.001	652 (72.2)	<0.001	240 (72.9)	0.005	130 (69.9)	0.239
Current (or stopped < 2y) smoking, n (%)	978 (23.1)	144 (14.6)	<0.001	112 (12.4)	<0.001	32 (9.7)	<0.001	59 (37.8)	<0.001
Hypercholesterolemia, n (%)	2282 (42.0)	508 (41.5)	0.773	375 (41.5)	0.913	133 (40.4)	0.686	85 (45.7)	0.327
Diabetes mellitus, n (%)	889 (16.4)	297 (24.2)	<0.001	211 (23.4)	<0.001	86 (26.1)	<0.001	30 (16.1)	1.000
Coronary artery disease, n (%)	953 (17.6)	338 (27.7)	<0.001	238 (26.4)	<0.001	100 (30.4)	<0.001	32 (17.2)	0.866
Prior stroke, n (%)	738 (13.6)	221 (18.1)	<0.001	161 (18)	<0.001	60 (18.2)	0.017	23 (12.4)	0.743
Poor outcome, n (%)	2151 (39.5)	684 (55.5)	<0.001	478 (52.9)	<0.001	206 (62.6)	<0.001	67 (36.0)	0.360
Mortality, n (%)	613 (11.3)	276 (22.4)	<0.001	187 (20.7)	<0.001	89 (27.1)	<0.001	19 (10.2)	0.724
Symptomatic ICH (ECASS-II criteria), n (%)	235 (4.3)	56 (4.5)	0.700	43 (4.8)	0.539	13 (4)	0.889	6 (3.2)	0.582
Recurrent stroke ^f	67 (3.4)	17 (3.3)	0.883	13 (3.6)	0.905	4 (2.7)	0.6213 (4.9)	0.538	

ECASS-II-criteria: European Cooperative Acute Stroke Study II-criteria; eGFR: estimated glomerular filtration rate; ICH: intracerebral haemorrhage; IQR: interquartile range; IVT: intravenous thrombolysis; NIHSS: National Institutes of Health Stroke Scale.

^aAnaemia = haemoglobin female: <12 g/dl; male: <13 g/dl.

^bMild anaemia = haemoglobin female: 11–11.9 g/dl; male: 11–12.9 g/dl.

^cModerate/severe anaemia = haemoglobin <11 g/dl.

^dPolyglobulia = haemoglobin female: >15.5 g/dl; male: >17 g/dl.

^eType of antithrombotic treatment not known in 152 patients.

^fInformation available in a subgroup of patients only, n = 2518.

Table 2. Univariate analysis of clinical characteristics (odds ratio with 95% confidence interval) in IVT patients.

Putative predicting variables	sICH	Poor outcome ^a	Mortality
Age (each year)	1.01 (1.01–1.03) $p < 0.001$	1.05 (1.04–1.05) $p < 0.001$	1.07 (1.06–1.08) $p < 0.001$
Sex	1.07 (0.88–1.23) $p = 0.516$	1.51 (1.37–1.67) $p < 0.001$	1.34 (1.16–1.54) $p < 0.001$
NIHSS (each point)	1.07 (1.06–1.09) $p < 0.001$	1.18 (1.17–1.20) $p < 0.001$	1.16 (1.14–1.17) $p < 0.001$
Systolic blood pressure (each mmHg)	1.00 (1.00–1.01) $p = 0.032$	1.00 (1.00–1.01) $p = 0.011$	1.00 (1.00–1.01) $p = 0.032$
Stroke-to-needle (each minute)	1.00 (0.99–1.00) $p = 0.235$	1.00 (1.00–1.00) $p = 0.136$	1.00 (0.99–1.00) $p = 0.389$
Decreasing eGFR (by 10 ml/min)	1.00 (1.00–1.01) $p < 0.001$	1.02 (1.02–1.02) $p < 0.001$	1.03 (1.02–1.03) $p < 0.001$
Glucose (each mmol/l)	1.03 (1.01–1.06) $p = 0.004$	1.09 (1.07–1.11) $p < 0.001$	1.09 (1.06–1.11) $p < 0.001$
Prior antithrombotics	1.22 (0.96–1.55) $p = 0.113$	1.47 (1.33–1.63) $p < 0.001$	1.99 (1.73–2.29) $p < 0.001$
Antiplatelets	1.20 (0.93–1.55) $p = 0.163$	1.34 (1.20–1.40) $p < 0.001$	1.71 (1.47–1.99) $p < 0.001$
Anticoagulation with or without antiplatelets	0.93 (0.50–1.72) $p = 0.817$	1.88 (1.47–2.41) $p < 0.001$	2.12 (1.59–2.83) $p < 0.001$
Atrial fibrillation	1.45 (1.19–1.78) $p < 0.001$	1.93 (1.74–2.16) $p < 0.001$	2.01 (1.74–2.33) $p < 0.001$
Hypertension	1.19 (0.96–1.46) $p = 0.107$	1.50 (1.35–1.66) $p < 0.001$	1.61 (1.37–1.89) $p < 0.001$
Smoking	0.67 (0.51–0.90) $p = 0.007$	0.70 (0.62–0.81) $p < 0.001$	0.50 (0.40–0.63) $p < 0.001$
Hypercholesterolemia	1.10 (0.91–1.33) $p = 0.321$	0.86 (0.78–0.95) $p = 0.003$	0.85 (0.74–0.98) $p = 0.025$
Diabetes mellitus	1.33 (1.06–1.68) $p = 0.014$	1.67 (1.47–1.87) $p < 0.001$	1.67 (1.41–1.97) $p < 0.001$
Coronary artery disease	1.38 (1.10–1.72) $p = 0.005$	1.44 (1.27–1.62) $p < 0.001$	1.95 (1.67–2.28) $p < 0.001$
Prior stroke	1.17 (0.90–1.51) $p = 0.237$	1.35 (1.18–1.54) $p < 0.001$	1.48 (1.24–1.78) $p < 0.001$
Anaemia ^b versus normal HL	1.04 (0.82–1.33) $p = 0.744$	1.91 (1.69–2.17) $p < 0.001$	2.28 (1.94–2.67) $p < 0.001$
Mild anaemia ^c versus normal HL	1.11 (0.80–1.55) $p = 0.542$	1.72 (1.50–1.99) $p < 0.001$	2.06 (1.72–2.47) $p < 0.001$
Moderate/severe anaemia ^d versus normal HL	0.91 (0.52–1.61) $p = 0.753$	2.57 (2.04–3.23) $p < 0.001$	2.93 (2.26–3.78) $p < 0.001$
Polyglobulia ^e versus normal HL	0.79 (0.41–1.50) $p = 0.469$	0.86 (0.64–1.17) $p = 0.343$	0.90 (0.55–1.45) $p = 0.660$
Decreasing HL (by 1 g/dl)	1.04 (0.98–1.10) $p = 0.160$	1.22 (1.19–1.26) $p < 0.001$	1.24 (1.19–1.29) $p < 0.001$

eGFR: estimated glomerular filtration rate; HL: haemoglobin level; IVT: intravenous thrombolysis; NIHSS: National Institutes of Health Stroke Scale; sICH: symptomatic intracerebral haemorrhage (ECASS II definition).

^aPoor outcome: modified Rankin Scale 3–6.

^bAnaemia = haemoglobin female: < 12 g/dl; male: < 13 g/dl.

^cMild anaemia = haemoglobin female: 11–11.9 g/dl; male: 11–12.9 g/dl.

^dModerate/severe anaemia = haemoglobin < 11 g/dl.

^ePolyglobulia = haemoglobin female: > 15.5 g/dl; male: > 17 g/dl.

(OR_{unadjusted} 2.28, 95% CI 1.94–2.67 and OR_{adjusted} 1.58, 95% CI 1.27–1.95) but not with sICH (OR_{unadjusted} 1.04, 95% CI 0.82–1.33 and OR_{adjusted} 0.94, 95% CI 0.69–1.30). The lack of association between HL and sICH remained irrespective of type of antithrombotic treatment (antiplatelet and/or anticoagulation) (Tables 2 and 3).

Mild and moderate to severe anaemia versus normal HL

Of 1232 patients with anaemia, 903 (73.3%) had mild anaemia and 329 (26.7%) moderate/severe anaemia. Baseline characteristics of both anaemia subgroups are presented in Table 1. Patients with mild and moderate/severe anaemia had more frequently poor functional outcome (52.9 and 62.6% versus 39.5% in patients with normal HL) and mortality (20.7 and 27.1% versus 11.3% in normal HL). Frequency of sICH did not differ significantly between patients with mild anaemia (4.8%), moderate/severe anaemia

(4.0%) and normal HL (4.3%) (Table 1). Mild anaemia was associated with poor functional outcome (OR_{unadjusted} 1.72, 95% CI 1.50–1.99) and mortality (OR_{unadjusted} 2.06, 95% CI 1.72–2.47). These associations remained significant after adjustment for potential confounders (poor functional outcome: OR 1.29, 95% CI 1.07–1.55 and mortality: OR 1.45, 95% CI 1.15–1.84). Similarly, moderate/severe anaemia was associated with poor functional outcome (OR_{unadjusted} 2.57, 95% CI 2.04–3.23; OR_{adjusted} 1.48, 95% CI 1.09–2.02) and mortality (OR_{unadjusted} 2.93, 95% CI 2.26–3.78; OR_{adjusted} 2.0, 95% CI 1.46–2.75). Neither mild nor moderate/severe anaemia was associated with the occurrence of sICH (Tables 2 and 3).

Polyglobulia versus normal HL

Compared to normal HL, polyglobulia ($n = 186$; 2.71%) did not significantly change the odds for any outcome (poor functional outcome OR_{unadjusted} 0.86, 95% CI 0.64–1.17; mortality OR_{unadjusted} 0.9, 95% CI

Table 3. Multivariate analysis of outcomes (odds adjusted for variables with $p < 0.05$ in the univariate analysis). Odds ratio (95% confidence interval), p -value.

Putative predicting variables	Outcome measures		
	sICH	Poor outcome ^a	Mortality
Anaemia versus normal HL	0.94 (0.69–1.30) ^b $p = 0.718$	1.25 (1.05–1.48) ^c $p = 0.012$	1.58 (1.27–1.95) ^d $p < 0.001$
Mild anaemia versus normal HL	1.03 (0.73–1.46) ^e $p = 0.865$	1.29 (1.07–1.55) ^f $p = 0.009$	1.45 (1.15–1.84) ^g $p = 0.002$
Moderate/severe anaemia versus normal Hb	0.72 (0.40–1.30) ^h $p = 0.277$	1.48 (1.09–2.02) ⁱ $p = 0.013$	2.00 (1.46–2.75) ^j $p < 0.001$
Decreasing HL (by 1 g/dl)	1.02 (0.94–1.11) ^k $p = 0.652$	1.07 (1.02–1.11) ^l $p = 0.004$	1.08 (1.02–1.15) ^m $p = 0.010$

HL: haemoglobin level; sICH: symptomatic intracerebral haemorrhage (ECASS II definition).

^aPoor outcome: modified Rankin Scale 3–6.

^bNIHSS on admission, age, CKD-EPI, pre-antithrombotic any, anaemia.

^cNIHSS on admission, age, glucose on admission, CKD-EPI, RF diabetes, RF hypertension, RF coronary artery disease, gender, smoking, pre-antithrombotic any, RF atrial fibrillation, RF hypercholesterolemia, RR systole on admission, RF prior ischaemic stroke, anaemia.

^dNIHSS on admission, age, glucose on admission, CKD-EPI, RF diabetes, RF hypertension, RF coronary artery disease, gender, smoking, pre-antithrombotic any, RF atrial fibrillation, RF hypercholesterolemia, RR systole on admission, RF prior ischaemic stroke, anaemia.

^eNIHSS on admission, age, CKD-EPI, pre-antithrombotic any, mild anaemia.

^fNIHSS on admission, age, stroke-to-needle-time, glucose on admission, CKD-EPI, RF diabetes, RF hypertension, RF coronary artery disease, gender, pre-antithrombotic any, RF atrial fibrillation, RF hypercholesterolemia, RR systole on admission, RF prior ischaemic stroke, mild anaemia.

^gNIHSS on admission, age, stroke-to-needle-time, glucose on admission, CKD-EPI, RF diabetes, RF hypertension, RF coronary artery disease, gender, pre-antithrombotic any, RF atrial fibrillation, RF hypercholesterolemia, RR systole on admission, RF prior ischaemic stroke, mild anaemia.

^hNIHSS on admission, severe anaemia.

ⁱNIHSS on admission, age, stroke-to-needle-time, glucose on admission, CKD-EPI, RF diabetes, RF hypertension, RF coronary artery disease, gender, pre-antithrombotic any, RF atrial fibrillation, RF hypercholesterolemia, RR systole on admission, RF prior ischaemic stroke, current smoking, severe anaemia.

^jNIHSS on admission, age, gender, glucose on admission, CKD-EPI, prior stroke, RR on admission, severe anaemia.

^kNIHSS on admission, age, glucose on admission, CKD-EPI, RF diabetes, RF coronary artery disease, smoking, pre-antithrombotic any, RF atrial fibrillation, RR systole on admission, decreasing HL in g/dl.

^lNIHSS on admission, age, glucose on admission, CKD-EPI, RF diabetes, RF hypertension, RF coronary artery disease, gender, smoking, pre-antithrombotic any, RF atrial fibrillation, RF hypercholesterolemia, RR systole on admission, RF prior ischaemic stroke, decreasing HL in g/dl.

^mNIHSS on admission, age, glucose on admission, CKD-EPI, RF diabetes, RF hypertension, RF coronary artery disease, gender, smoking, pre-antithrombotic any, RF atrial fibrillation, RF hypercholesterolemia, RR systole on admission, RF prior ischaemic stroke, decreasing HL in g/dl.

0.55–1.45; sICH OR_{unadjusted} 0.79, 95% CI 0.41–1.50) (Tables 1 to 3).

HL as a continuous variable

The adjusted logistic regression model showed an increasing probability of poor outcome and mortality with decreasing HL. A decrease in HL by 1 g/dl was associated with an OR of 1.07 (95% CI 1.02–1.11) for poor outcome and with an OR of 1.08 (95% CI 1.02–1.15) for mortality. HLs were not significantly associated with sICH (Figure 1 and Table 3).

The ability of HL to predict poor functional outcome or mortality was low (AUC of ROC curve for poor functional outcome: 0.59 (95% CI 0.58–0.61, $p < 0.001$); for mortality: 0.60 (95% CI 0.58–0.62, $p < 0.001$) (eFigure 1).

Discussion

This study showed the following key results for the association between HL and outcomes in acute ischaemic stroke patients treated with IVT: (i) 17.9% of

IVT-treated stroke patients had anaemia on admission. (ii) Anaemia was an independent predictor for poor functional outcome and mortality in IVT-treated stroke patients, but not for symptomatic ICH. (iii) The more severe the anaemia the higher the probability of poor functional outcome and mortality. (iv) Polyglobulia was not associated with any outcome.

In our study population, 17.9% of patients had anaemia (according to the WHO criteria) on admission and before IVT administration. Presence of anaemia independently increased the probability of poor functional outcome after three months of follow-up by 25% and the probability of mortality by 60%. Previously, only one smaller observational study ($n = 217$) has investigated the impact of HL on outcomes in IVT-treated stroke patients.⁷ Although the proportion of patients with anaemia on admission was higher in patients with poor functional outcome (20.4%) compared to patients with favourable functional outcome (10.5%; $p = 0.04$), HLs on admission were not independently associated with poor functional outcome ($p = 0.20$) or mortality ($p = 0.45$). However, the

number of patients with anaemia was small ($n=33$), though the proportion (i.e. 15%) resembled that of the present study (i.e. 17.9%). Interestingly, after inclusion of patients who had developed anaemia during the first five days of hospitalisation into analysis ($n=86$), anaemia was independently associated with poor functional outcome (OR 2.61, 95% CI 1.33–5.11) but still not with mortality ($p=0.13$). A recent study, including IVT and endovascular treated patients, found that anaemia at hospital admission and any decrease of haemoglobin were associated with poor functional outcome and higher mortality.⁸

The large sample size in the present study allowed a subgroup analysis of patients with mild and moderate/severe anaemia. Data on anaemia subgroups are scarce. One retrospective study after mechanical thrombectomy²³ ($n=90$) reported an association between severe anaemic patients ($Hb<10$ g/dl) and poor functional outcome but not for patients with mild anaemia. In our study, both mild and moderate/severe anaemic patients were independently associated with poor functional outcome and mortality but the odds were higher in the moderate/severe group. In line, the probability for poor functional outcome and mortality increased with decreasing HL. Every decrease in HL by 1 g/dl increased the probability of poor outcome by 7% and the probability of mortality by 8%.

Several underlying pathophysiological mechanisms could explain the association between anaemia and an increased probability of poor functional outcome and mortality. Arterial oxygen content depends on HL and arterial oxygen saturation. The oxygen content and cerebral blood flow determine oxygen delivery to the brain. Consequently, decreasing HL lowers the arterial oxygen content and possibly leads to changes in blood flow resulting in an impaired tissue oxygen supply. Therefore, hypoxia in penumbral lesions may be increased leading to more extensive ischaemic areas.^{24,25}

Furthermore, anaemia could compromise the cerebral autoregulation which maintains cerebral blood flow and oxygen carrying capacity through collaterals to penumbral lesions.^{26,27} Healthy patients with anaemia might tolerate hypoxia with an HL of approximately down to 8 g/dl.²⁵ However, in anaemic stroke patients who likely have additional comorbidities, cerebral autoregulation is thought to be already impaired.²⁸ In addition, a mathematical model of regional cerebral oxygen uptake also suggested that oxygen uptake in ischaemic penumbra progressively decreases below an HL of 10 g/dl.²⁹ Other potential pathophysiological mechanisms include the modulation of adhesion molecules by hyperdynamic circulation in anaemic patients³⁰ and the upregulation of inflammatory mediators.^{1,31} In line, a recent study found a positive

correlation between decreasing baseline HL and increasing final infarct volume in acute stroke patients.⁸

The clear and independent association between anaemia on admission and poor functional outcome and mortality in acute ischaemic stroke patients carries the chance for hyperacute interventions. Such an intervention could be red blood cell transfusion (RBCT). In one retrospective study investigating the optimal management of HL and RBCT in patients with severe ischaemic stroke, RBCT was not associated with any clinical improvement.³² However, the sample size was small ($n=109$), and stroke severity was high (median NIHSS 19). Furthermore, RBCT was performed at the discretion of the neurologic ICU physician in charge, based on the general aim to keep HL between 8 and 10 g/dl over the course of the hospitalisation and not necessarily during the hyperacute phase when the penumbra might still be preserved.

Furthermore, the optimal HL threshold for RBCT is unclear.³³ In general, RBCT is not considered in patients with $HL>10$ g/dl. In some ischaemic populations (i.e. acute coronary syndrome) the threshold for RBCT is between 8 and 10 g/dl.³⁴ Data on RBCT in ischaemic patients (i.e. myocardial infarction) with anaemia remain ambiguous: in some studies, RBCT reduced mortality^{35,36} while in others RBCT increased adverse outcomes after percutaneous coronary intervention³⁷ and in ST-elevation myocardial infarction.³⁸ In anaemic patients with aneurysmal subarachnoid haemorrhage, RBCTs are suggested to be beneficial when there was no considerable anaemia beforehand.³⁹ On the other hand, in the setting of perioperative procedures RBCT was suggested to be associated with stroke.⁴⁰ In patients undergoing coronary artery bypass surgery, the use of solvent/detergent treated plasma and platelet transfusions seemed to have a larger impact on the development of stroke than RBCT.⁴¹

Nevertheless, the strong association of anaemia with poor functional outcome and mortality in acute setting of IVT-treated stroke patients and the high vulnerability of penumbral brain tissue to hypoxia might justify a randomised controlled trial investigating RBCT in the hyperacute phase of ischaemic stroke.

In our study, polyglobulia was not associated with functional outcome, mortality or sICH. In the general stroke population, a U-shaped relationship between HL and short-term mortality (up to one month) in men has been suggested.¹ This might indicate that patients with polyglobulia have a special benefit of IVT. However, the proportion of patients with polyglobulia was small (2.7%; $n=186$) in our study and results should be interpreted with caution.

Our study has the following strengths: (i) the large sample size reduced the risk of chance findings and

allowed adjustment for several confounding variables, (ii) the low number of missing data on HL (2.9%) and clinical outcome at three months (3.5%) reduced the risk of bias and (iii) the prospective and systematic approach of data assessment was not influenced by the current research question.

This study has limitations: (i) the TRISP registry is not monitored and does not provide data from comparison group of patients not treated with IVT. Thus, we were not able to calculate the treatment effect of IVT stratified to HL categories. (ii) We were neither able to classify the type of anaemia nor did we have any information about the underlying cause of anaemia, which may affect functional three-month outcome and mortality, e.g. in the case of cancer-related anaemia, anaemia of chronic disease or malnutrition. Due to insufficient data it was also not possible to fully incorporate potential confounders such as frailty (i.e. by a standard measure of comorbidity) which is associated with anaemia and poor outcome. However, we were able to adjust for parameters that are relevant in anaemia of chronic disease most importantly for renal function (via estimated glomerular filtration rate) and for vascular risk factors. Hence chronic disease as a possible confounder on functional outcome and mortality is partly addressed in the multivariate analyses. In addition, anaemia might be a manifestation of chronic, e.g. gastrointestinal, bleeding. In these cases, IVT treatment is likely to deteriorate outcomes. However, because IVT is contraindicated in haemorrhagic diseases (acute or chronic), the number of these patients is expected to be very small. (iii) Further, we have no information on the course of the HL over time and whether HLs were deliberately altered with RBCT or iron supplements. (iv) The investigated data were collected from high volume stroke centres. These results may not apply to low volume centres.

Conclusions

Anaemia was independently associated with poor clinical outcome and mortality in IVT-treated stroke patients. The more severe the anaemia, the stronger the association with poor clinical outcome and mortality. Therefore, severe anaemia might be a target for interventions in hyperacute stroke. Anaemia was not associated with occurrence of sICH and no significant association between polyglobulia and any outcome was found.

Declaration of Conflicting Interests

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Ethical approval

The study was approved by the ethics committee in Basel, Switzerland.

Informed consent

The requirement for additional local ethical approval/informed consent differed between participating centres and was obtained if required. Anonymised data will be shared by request from any qualified investigator.

Guarantor

HG.


Contributorship

VLA designed/conceptualised the study, collected data, analysed/interpreted the data, drafted the manuscript. LK designed/conceptualised the study, collected data, revised the manuscript. ASA analysed/interpreted data and revised the manuscript. STE and HG designed/conceptualised and initiated the study, supervised the study, collected data, analysed/interpreted the data, revised the manuscript. All other authors collected data and revised the manuscript.


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